

NBS1/NBN Rabbit pAb

Catalog No.: A0782

Basic Information

Observed MW

105kDa

Calculated MW

85kDa

Category

Primary antibody

Applications

WB,FC

Cross-Reactivity

Human

Background

Mutations in this gene are associated with Nijmegen breakage syndrome, an autosomal recessive chromosomal instability syndrome characterized by microcephaly, growth retardation, immunodeficiency, and cancer predisposition. The encoded protein is a member of the MRE11/RAD50 double-strand break repair complex which consists of 5 proteins. This gene product is thought to be involved in DNA double-strand break repair and DNA damage-induced checkpoint activation.

Recommended Dilutions

WB	1:500 - 1:1000
FC	1:20 - 1:50

Immunogen Information

Gene ID	Swiss Prot
4683	O60934

Immunogen

A synthetic peptide of human NBS1/NBS1/NBN

Synonyms

ATV; NBS; P95; NBS1; AT-V1; AT-V2; NBS1/NBN

Contact

 | www.abclonal.com

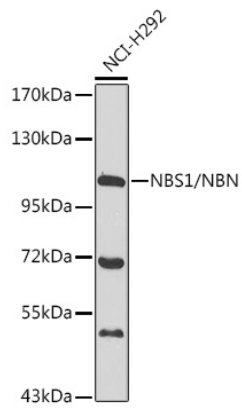
Product Information

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

Storage

Store at 4°C. Avoid freeze / thaw cycles.
Buffer: PBS with 0.02% sodium azide,pH7.3.

Validation Data



Western blot analysis of extracts of NCI-H292 cells, using NBS1/NBS1/NBN antibody (A0782).
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.
Lysates/proteins: 25 μ g per lane.
Blocking buffer: 3% nonfat dry milk in TBST.